

### **Remarks/Arguments**

After entry of this Amendment, claims 25, 26, and 28-30, as amended, will be pending in the application. Claims 1-24, 27, and 31-38 have been canceled without prejudice. The right to prosecute the subject matter of any of the canceled claims in this or in a continuation, continuation-in-part, or divisional application is hereby expressly reserved.

The undersigned thanks Examiner Choi for the recitation in the Advisory Action mailed June 7, 2010 that present claims 25, 26, and 28-30 “would be allowable if submitted in a separate, timely filed amendment cancelling the non-allowable claims.” The present Amendment cancels the alleged “non-allowable” claims without prejudice. The present Amendment also repeats those remarks and arguments found persuasive in the Amendment under 37 C.F.R. § 1.116 filed on May 17, 2010, which was not entered by the Office, in order to make those remarks and arguments of record.

#### **I. Interview Summary**

The undersigned appreciates the courtesies extended to her colleague Esther Kepplinger, Reg. No. 57,243, by Examiner Choi in the personal interview held on May 3, 2010. As stated in the Amendment under 37 C.F.R. § 1.116 filed on May 17, 2010, in the Interview, the rejections of the claims under 35 U.S.C. § 103(a) were discussed, and Examiner acknowledged that Applicant’s arguments and amendments would appear to overcome the rejections of record. Other topics discussed in the Interview are included in the Remarks that follow.

#### **II. Claim Amendments**

Claims 28-30 have been amended to depend from pending claim 25, rather than from canceled claim 1. No new matter has been added to the claims by these amendments.

#### **III. Claim Rejections under 35 U.S.C. § 112**

Claims 35-38 stand rejected under 35 U.S.C. § 112, first paragraph as allegedly failing to comply with the written description requirement for the recitation that “the composition does not comprise an amino acid.” This rejection has been rendered moot by the cancellation of claims 35-38 without prejudice.

Claims 28-30 stand rejected under 35 U.S.C. § 112, second paragraph as allegedly indefinite for being dependent on canceled claim 1. This rejection has been rendered moot by the amendment of claims 28-30 to depend from pending claim 25, rather than from canceled claim 1.

For these reasons, it is respectfully requested that the Office withdraw the claim rejections under 35 U.S.C. § 112.

IV. Claim Rejections under 35 U.S.C. § 103

Claims 25, 26, and 31-34 stand rejected under 35 U.S.C. § 103(a) as allegedly obvious over U.S. Patent No. 4,938,969 to Schinitzky and Meisner (“Schinitzky & Meisner”) in view of U.S. Patent No. 5,804,594 to Murad (“Murad”); U.S. Patent No. 5,902,591 to Herstein (“Herstein”) or U.S. Patent No. 5,140,043 to Darr and Pinnell (“Darr”); THE MERCK INDEX, entry 855 (9th ed. 1976) (“Merck Index”); and J.P. Yuan & F. Chen, *J. Agric. Food Chem.*, 46: 5078-82 (1998) (“Yuan”). This rejection has been rendered moot as to claims 31-34 by the cancellation of those claims without prejudice. Arguments in support of the patentability of claims 25 and 26, as well as claims 28-30, over these references are presented below.

Claims 25, 26, and 28-30 recite topical compositions comprising: at least 10% (w/v) ascorbic acid; approximately 10% to 25% (w/v) glucosamine; and water, wherein the composition has a pH of about 3.5 to about 4.1; and wherein the composition is prepared by a process comprising: (a) dissolving about 10% to about 50% of the ascorbic acid in water at a temperature of between about 60°C to about 90°C to provide an aqueous ascorbic acid solution of at least 20% (w/v); (b) cooling the aqueous ascorbic acid solution to below about 40°C; (c) combining the aqueous ascorbic acid solution with water, glucosamine, and ascorbic acid to provide a composition comprising water, approximately 10% to 25% (w/v) glucosamine and at least 10% (w/v) ascorbic acid; and (d) adjusting the pH of the composition to about 3.5 to about 4.1.

A. The references cited by the Office, even when combined, do not teach or suggest all of the recitations of claims 25, 26, and 28-30

1. “(a) dissolving about 10% to about 50% of the ascorbic acid in water at a temperature of between about 60°C to about 90°C to provide an aqueous

ascorbic acid solution of at least 20% (w/v); (b) cooling the aqueous ascorbic acid solution to below about 40°C”

As previously discussed, none of Schinitsky & Meisner, Murad, Herstein, or Darr teaches or suggests a composition comprising ascorbic acid that has been prepared by a process comprising the steps of “(a) dissolving about 10% to about 50% of the ascorbic acid in water at a temperature of between about 60°C to about 90°C to provide an aqueous ascorbic acid solution of at least 20% (w/v)” and “(b) cooling the aqueous ascorbic acid solution to below about 40°C,” as recited in claims 25, 26, and 28-30. (*See, e.g.*, Amendment filed December 3, 2009). The Office cites Merck Index and Yuan to provide the missing teaching of these process steps. This is unavailing for at least the following reasons.

Yuan teaches away from exposing ascorbic acid to a temperature of between about 60°C to about 90°C, as recited in the claims, by showing that heating ascorbic acid at 60°C causes the ascorbic acid to degrade. For example, Yuan reports the presence of at least three degradation products after heating a solution of ascorbic acid in aqueous media at 60°C. (Yuan, pp. 5079, 5081-82, Figure 1). The Office has flatly dismissed this argument, stating: “The Yuan et al. reference does not teach away from the claimed invention as Yuan et al. discloses that there is little degradation at 60 degrees compared to 100 degrees Celsius.” (Office Action, p. 10). No matter whether Yuan teaches that there is less degradation at 60°C than at 100°C, Yuan teaches degradation at 60°C and, therefore, one of ordinary skill in the art would be motivated by Yuan as a whole to avoid heating at 60°C in order to avoid any degradation of the ascorbic acid.

Further, Merck Index merely discloses the solubility of ascorbic acid in water (80% at 100°C and 40% at 45°C) and does not and would not teach or suggest to the person of ordinary skill in the art a topical composition comprising ascorbic acid, or a method for preparing such a topical composition that includes steps (a) and (b) above. From the Merck Index, one would expect the solubility to decrease with cooling and thus provides no motivation for heating in the first place.

Therefore, neither Merck Index nor Yuan, when combined with the teachings of Schinitsky & Meisner, Murad, Herstein and/or Darr, teaches or suggests a topical composition of ascorbic acid made by a process including steps (a) and (b) above, as recited in the claims.

2. “wherein the composition has a pH of about 3.5 to about 4.1”

As previously discussed, none of Schinitzky & Meisner, Murad, Merck Index or Yuan teaches or suggests a composition comprising ascorbic acid that has a pH of about 3.5 to about 4.1, as recited in claims 25, 26, and 28-30. (*See, e.g.*, Amendment filed December 3, 2009). The Office cites Herstein or Darr to provide the missing teaching of the recited pH. This is unavailing for at least the following reasons.

The Declaration of Dr. Lorraine Faxon Meisner under 37 C.F.R. § 1.132 submitted on December 3, 2009 (“Meisner Declaration”) clearly establishes through several peer-reviewed journal articles that, at the time of the invention, it would have been entirely unexpected that an ascorbic acid composition would be stable enough for use in a topical composition at a pH of about 3.5 to about 4.1. (Meisner Declaration, ¶¶ 11-12). Dr. Meisner’s conclusion is supported not only by the Bauernfeind, Hajratwala, and Kassem articles cited in her Declaration, but also by Herstein and Darr, which were cited by the Office.

Herstein and Darr both acknowledge the instability of ascorbic acid compositions at the pH range recited in the claims. For example, Herstein teaches that emulsions of ascorbic acid having a pH within the range of 3.5 to 4.1 are unstable if they lack an organoclay stabilizer: “As can be seen from the data, the physical appearance of the emulsion initially is acceptable and remains acceptable (no breaking of the emulsion) after 25 days. Without the organoclay ingredient, the emulsion would begin to break down after a few days, i.e., 2-3 days.” (Herstein, col. 13, ll. 35-41). In addition, Darr stresses the importance of maintaining the pH of an ascorbic acid composition at “no more than about 3 to 3.5, preferably no more than about 2.5” in order to “ensure[] that greater than 82% of the ascorbic acid remains in the protonated, uncharged form,” which “removes the ionic repulsion of the two oxygen groups, thus stabilizing the molecule.” (Darr, col. 3, ll. 29-30, col. 4, ll. 7-18).

Therefore, neither Herstein nor Darr, when combined with the teachings of Schinitzky & Meisner, Murad, Merck Index and/or Yuan, teaches or suggests the recited compositions having a pH of about 3.5 to about 4.1.

3. The ascorbic acid in the claimed compositions is surprisingly and unexpectedly more stable than native ascorbic acid

Moreover, as discussed in the Interview, Applicant’s Specification asserts that ascorbic acid that has been prepared by the process recited in the claims produces a topical composition

that is surprisingly and unexpectedly more stable than a topical composition having "native" ascorbic acid:

As an example, containers having a 1 to 20% (w/v) concentration of a mixture of pretreated ascorbic acid in a 1:1 to 1:10 ratio, together with ascorbic acid formulated under more standard conditions (i.e., dissolved or added in solid form to a formulation at temperatures of about 20 to about 40°C--"native ascorbic acid") were quite stable when shipped and/or stored under adverse conditions, or even when heated. **The stability of such formulations was enhanced in comparison to conventional low pH formulations containing untreated ascorbic acid**, e.g., low pH creams containing 10% (w/v) untreated ascorbic acid. It is postulated that the observed stability of the present compositions is afforded by an equilibrium reaction between ascorbic acid and monhydroascorbic acid that maintains a stable solution of ascorbic acid.

(Specification, page 13, ¶ 30).

For these reasons, the Office has failed to make out a *prima facie* case of obviousness of any of claims 25, 26 or 28-30. Accordingly, the rejections of claims 25 and 26 under 35 U.S.C. § 103 as obvious over Schinitsky & Meisner in view of Murad, Herstein or Darr, Merck Index, and Yuan cannot stand and should be withdrawn. Further, any rejection of claims 28-30 under 35 U.S.C. § 103(a) as obvious over the cited references is obviated.

#### V. Conclusion

In view of the foregoing amendments and remarks, it is respectfully submitted that the claims are in condition for allowance. Early and favorable action by the Examiner is earnestly solicited. If any outstanding issues remain, the Examiner is invited to contact the undersigned at (212) 497-7731 to discuss the same.

Application No. 09/997,663  
Amdt. filed July 19, 2010  
Reply to Office Action mailed February 17, 2010

No fee is believed to be due for the submission of this response. Should any fees be required, please charge all such fees to Wilson, Sonsini, Goodrich & Rosati Deposit Account No. 23-2415 (36091-701.501).

Respectfully submitted,

Dated: July 19, 2010

By: /Gina R. Gencarelli/  
Gina R. Gencarelli  
Reg. No. 59,729

WILSON, SONSINI, GOODRICH & ROSATI PC  
650 Page Mill Road  
Palo Alto, CA 94304  
Phone: (650) 493-9300  
Fax: (650) 493-6811  
**Customer No. 21971**